

PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details:
<http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf>

=>
Uploading 09980593.str

L1 STRUCTURE UPLOADED

=> d 11
L1 HAS NO ANSWERS
L1 STR
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

Structure attributes must be viewed using STN Express query preparation.

=> s 11
SAMPLE SEARCH INITIATED 09:32:16 FILE 'REGISTRY'
SAMPLE SCREEN SEARCH COMPLETED - 4520 TO ITERATE

22.1% PROCESSED 1000 ITERATIONS 0 ANSWERS
INCOMPLETE SEARCH (SYSTEM LIMIT EXCEEDED)
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS: ONLINE **COMPLETE**
BATCH **COMPLETE**
PROJECTED ITERATIONS: 86370 TO 94430
PROJECTED ANSWERS: 0 TO 0

L2 0 SEA SSS SAM L1

=> s 11 sss full
FULL SEARCH INITIATED 09:32:35 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 91344 TO ITERATE

100.0% PROCESSED 91344 ITERATIONS 0 ANSWERS
SEARCH TIME: 00.00.03

L3 0 SEA SSS FUL L1

=> log y
COST IN U.S. DOLLARS SINCE FILE TOTAL
ENTRY SESSION
FULL ESTIMATED COST 148.55 148.76

STN INTERNATIONAL LOGOFF AT 09:32:43 ON 19 MAR 2003

Broad
Search

Crossover limits have been increased. See HELP CROSSOVER for details.

Experimental and calculated property data are now available. See HELP PROPERTIES for more information. See STNote 27, Searching Properties in the CAS Registry File, for complete details:
<http://www.cas.org/ONLINE/STN/STNOTES/stnotes27.pdf>

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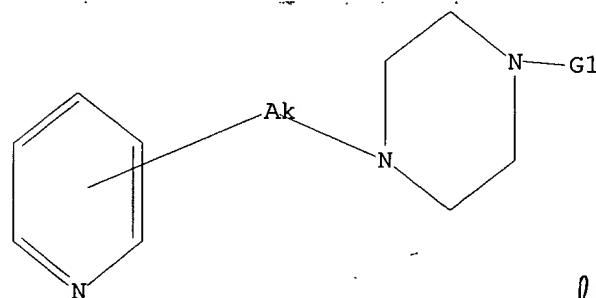
Uploading 09980593.str

L1 STRUCTURE UPLOADED

=> d 11

L1 HAS NO ANSWERS

L1 STR



G1 C, SO2

G2 C, N

*search also submitted
to library*

Structure attributes must

reparation.

=> s 11

SAMPLE SEARCH INITIATED 09:40:44
SAMPLE SCREEN SEARCH COMPI

15.2% PROCESSED 1000 ITERATIONS
INCOMPLETE SEARCH (SYSTEM 1)
SEARCH TIME: 00.00.01

2 ANSWERS

FULL FILE PROJECTIONS: ONLY

BATCH COMPLETE**

PROJECTED ITERATIONS: 126681 TO 136399

PROJECTED ANSWERS: 3339 TO 5079

L2 32 SEA SSS SAM L1

=> s 11 sss full

FULL SEARCH INITIATED 09:40:44 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED - 131073 TO ITERATE

100.0% PROCESSED 131073 ITERATIONS
SEARCH TIME: 00.00.03

3286 ANSWERS

L3 3286 SEA SSS FUL L1

=> file caplus		SINCE FILE	TOTAL
COST IN U.S. DOLLARS		ENTRY	SESSION
FULL ESTIMATED COST		148.15	148.36

FILE 'CAPLUS' ENTERED AT 09:40:54 ON 19 MAR 2003
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 PLEASE SEE "HELP USAGETERMS" FOR DETAILS.
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FILE COVERS 1907 - 19 Mar 2003 VOL 138 ISS 12
 FILE LAST UPDATED: 18 Mar 2003 (20030318/ED)

This file contains CAS Registry Numbers for easy and accurate substance identification.

=> s 13
 L4 1884 L3

=> s 14 and metalloprotein?
 L5 15 L4 AND METALLOPROTEIN?

=> d ibib abs hitstr tot

L5 ANSWER 1 OF 15 CAPLUS COPYRIGHT 2003 ACS
 ACCESSION NUMBER: 2003:5956 CAPLUS
 DOCUMENT NUMBER: 138:73254
 TITLE: Preparation of thiazolylaminopyridines as tyrosine kinase inhibitors with therapeutic uses
 INVENTOR(S): Bilodeau, Mark T.; Hartman, George D.
 PATENT ASSIGNEE(S): Merck & Co., Inc., USA
 SOURCE: PCT Int. Appl., 93 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003000687	A1	20030103	WO 2002-US21110	20020618
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,			

GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
 RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

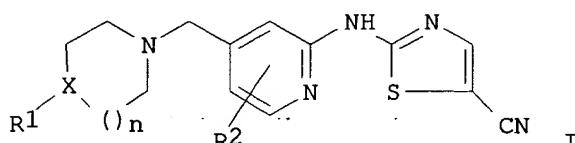
PRIORITY APPLN. INFO.:

US 2001-300245P P 20010622

OTHER SOURCE(S):

MARPAT 138:73254

GI



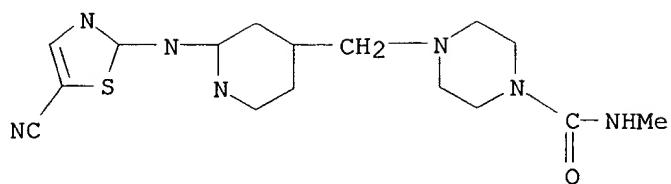
AB The present invention relates to thiazolylaminopyridines (shown as I; variables defined below; e.g. 4-[2-(5-cyanothiazol-2-ylamino)pyridin-4-ylmethyl]piperazine-1-carboxylic acid methylamide) which inhibit, regulate and/or modulate tyrosine kinase signal transduction, compns. which contain these compds., and methods of using them to treat tyrosine kinase-dependent diseases and conditions, such as angiogenesis, cancer, tumor growth, atherosclerosis, age related macular degeneration, diabetic retinopathy, inflammatory diseases, and the like in mammals. For I: n is 0 or 1; X is C-H or N, provided X is C-H if n = 1 and R1 is SO2-(C1-C6 alkyl) and provided that X is C-H if R1 is NH(C:O)NR3H; R1 is SO2(C1-C6 alkyl), (C:O)NR3H, or NH(C:O)NR3H; R2 is H, OH, OC1-C6 alkyl, C1-C6 alkyl, or halo; and R3 is C1-C6 alkyl. Compds. I inhibit VEGF-stimulated mitogenesis of human vascular endothelial cells in culture with IC50 values = 0.01-5.0 .mu.M. 4-[2-(5-Cyanothiazol-2-ylamino)pyridin-4-ylmethyl]piperazine-1-carboxylic acid methylamide, 2-[4-[(4-(methyldisulfonyl)piperidin-1-yl)methyl]pyridin-2-yl]amino]-1,3-thiazole-5-carbonitrile, and 4-[2-(5-cyanothiazol-2-ylamino)-3-methylpyridin-4-ylmethyl]piperazine-1-carboxylic acid methylamide show enhanced pharmacokinetic properties as compared to previously reported thiazolylaminopyridines in WO 01/17995 A1. Although the methods of prepn. are not claimed, 13 example prepn. are included.

IT **479611-82-0P**, 4-[2-(5-Cyanothiazol-2-ylamino)pyridin-4-ylmethyl]piperazine-1-carboxylic acid methylamide **479612-56-1P**, 4-[2-(5-Cyanothiazol-2-ylamino)-3-methylpyridin-4-ylmethyl]piperazine-1-carboxylic acid methylamide trifluoroacetate
 RL: PAC (Pharmacological activity); PKT (Pharmacokinetics); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; prepn. of thiazolylaminopyridines as tyrosine kinase inhibitors with therapeutic uses)

RN 479611-82-0 CAPLUS

CN 1-Piperazinecarboxamide, 4-[(2-(5-cyano-2-thiazolyl)amino)-4-pyridinyl]methyl-N-methyl- (9CI) (CA INDEX NAME)



*** FRAGMENT DIAGRAM IS INCOMPLETE ***

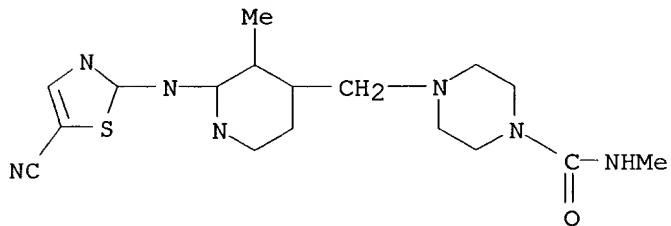
RN 479612-56-1 CAPLUS

CN 1-Piperazinecarboxamide, 4-[(2-[(5-cyano-2-thiazolyl)amino]-3-methyl-4-pyridinyl)methyl]-N-methyl-, trifluoroacetate (9CI) (CA INDEX NAME)

CM 1

CRN 479612-55-0

CMF C17 H21 N7 O S

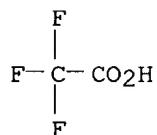


*** FRAGMENT DIAGRAM IS INCOMPLETE ***

CM 2

CRN 76-05-1

CMF C2 H F3 O2



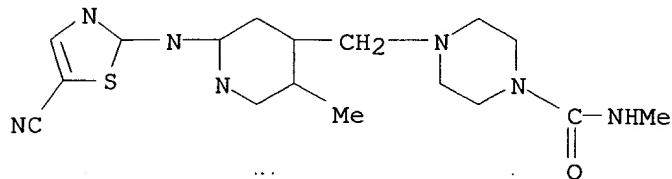
IT 479612-28-7P, 4-[2-(5-Cyanothiazol-2-ylamino)-5-methylpyridin-4-ylmethyl]piperazine-1-carboxylic acid methylamide 479612-29-8P, 4-[(2-(5-Cyanothiazol-2-ylamino)-5-methylpyridin-4-yl)methyl]piperazine-1-carboxylic acid methylamide trifluoroacetate 479612-55-0P, 4-[2-(5-Cyanothiazol-2-ylamino)-3-methylpyridin-4-ylmethyl]piperazine-1-carboxylic acid methylamide 479612-74-3P, 4-[[2-Chloro-6-[(5-cyano-1,3-thiazol-2-yl)amino]pyridin-4-yl)methyl]-N-methylpiperazine-1-carboxamide 479612-92-5P, 4-[[2-(5-Cyano-1,3-thiazol-2-yl)amino]-6-ethylpyridin-4-yl)methyl]-N-methylpiperazine-1-carboxamide 479613-12-2P, 2-[[4-[(4-Acetylpirazin-1-yl)methyl]-6-methylpyridin-2-yl]amino]-1,3-thiazole-5-carbonitrile 479613-13-3P, 2-[[4-[(4-Acetylpirazin-1-yl)methyl]-6-methylpyridin-2-yl]amino]-1,3-thiazole-5-carbonitrile trifluoroacetate

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(drug candidate; prepn. of thiazolylaminopyridines as tyrosine kinase inhibitors with therapeutic uses)

RN 479612-28-7 CAPLUS

CN 1-Piperazinecarboxamide, 4-[[2-[(5-cyano-2-thiazolyl)amino]-5-methyl-4-pyridinyl]methyl]-N-methyl- (9CI) (CA INDEX NAME)



*** FRAGMENT DIAGRAM IS INCOMPLETE ***

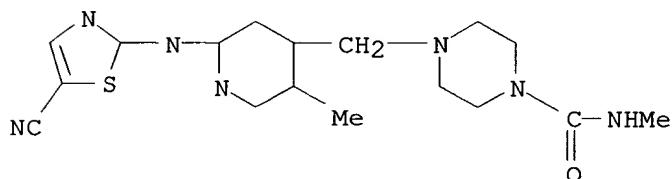
RN 479612-29-8 CAPLUS

CN 1-Piperazinecarboxamide, 4-[[2-[(5-cyano-2-thiazolyl)amino]-5-methyl-4-pyridinyl]methyl]-N-methyl-, trifluoroacetate (9CI) (CA INDEX NAME)

CM 1

CRN 479612-28-7

CMF C17 H21 N7 O S

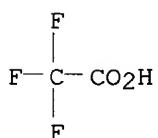


*** FRAGMENT DIAGRAM IS INCOMPLETE ***

CM 2

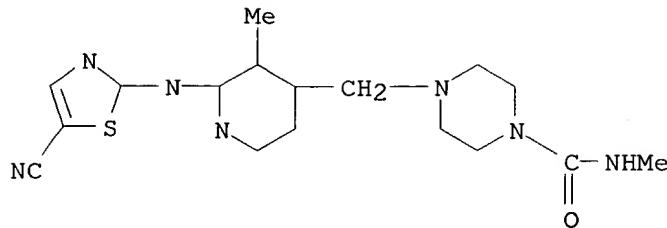
CRN 76-05-1

CMF C2 H F3 O2



RN 479612-55-0 CAPLUS

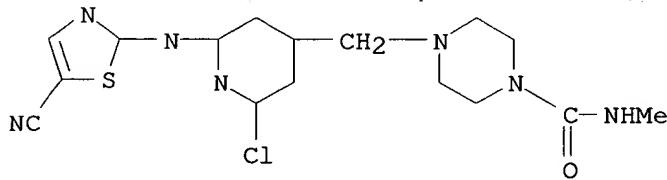
CN 1-Piperazinecarboxamide, 4-[[2-[(5-cyano-2-thiazolyl)amino]-3-methyl-4-pyridinyl]methyl]-N-methyl- (9CI) (CA INDEX NAME)



*** FRAGMENT DIAGRAM IS INCOMPLETE ***

RN 479612-74-3 CAPLUS

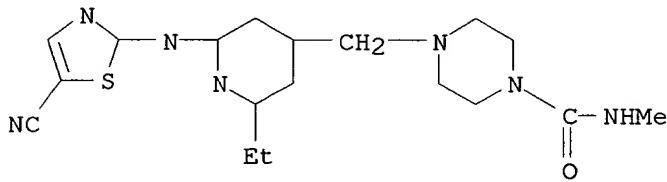
CN 1-Piperazinecarboxamide, 4-[(2-chloro-6-[(5-cyano-2-thiazolyl)amino]-4-pyridinyl)methyl]-N-methyl- (9CI) (CA INDEX NAME)



*** FRAGMENT DIAGRAM IS INCOMPLETE ***

RN 479612-92-5 CAPLUS

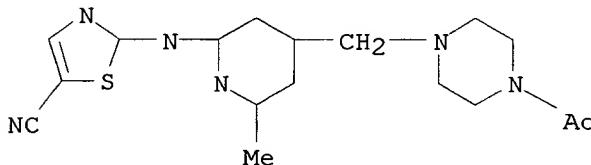
CN 1-Piperazinecarboxamide, 4-[(2-[(5-cyano-2-thiazolyl)amino]-6-ethyl-4-pyridinyl)methyl]-N-methyl- (9CI) (CA INDEX NAME)



*** FRAGMENT DIAGRAM IS INCOMPLETE ***

RN 479613-12-2 CAPLUS

CN Piperazine, 1-acetyl-4-[(2-[(5-cyano-2-thiazolyl)amino]-6-methyl-4-pyridinyl)methyl]- (9CI) (CA INDEX NAME)

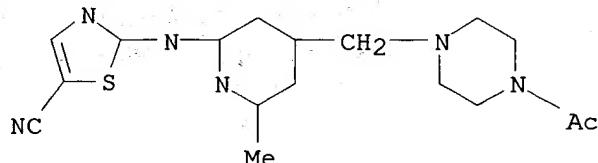


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RN 479613-13-3 CAPLUS

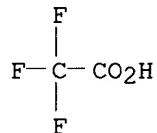
CN Piperazine, 1-acetyl-4-[(2-[(5-cyano-2-thiazolyl)amino]-6-methyl-4-pyridinyl)methyl]-, mono(trifluoroacetate) (9CI) (CA INDEX NAME)

CM 1

CRN 479613-12-2
CMF C17 H20 N6 O S

*** FRAGMENT DIAGRAM IS INCOMPLETE ***

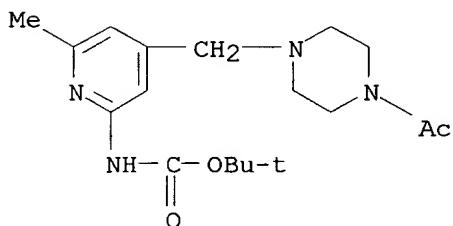
CM 2

CRN 76-05-1
CMF C2 H F3 O2

IT 479613-21-3P, tert-Butyl 4-[(4-acetyl-1-piperazinyl)methyl]-6-methylpyridin-2-ylcarbamate 479613-27-9P, 1-Acetyl-4-[(2-amino-6-methylpyridin-4-yl)methyl]piperazin-4-ium chloride
 RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
 (prepn. of thiazolylaminopyridines as tyrosine kinase inhibitors with therapeutic uses)

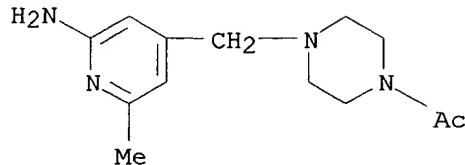
RN 479613-21-3 CAPLUS

CN Carbamic acid, [4-[(4-acetyl-1-piperazinyl)methyl]-6-methyl-2-pyridinyl]-, 1,1-dimethylethyl ester (9CI) (CA INDEX NAME)



RN 479613-27-9 CAPLUS

CN Piperazine, 1-acetyl-4-[(2-amino-6-methyl-4-pyridinyl)methyl]-, monohydrochloride (9CI) (CA INDEX NAME)



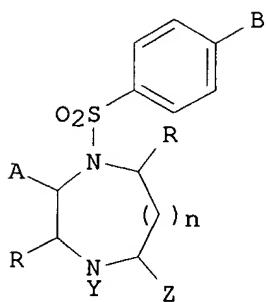
● HCl

REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 2 OF 15 CAPLUS COPYRIGHT 2003 ACS
ACCESSION NUMBER: 2002:849621 CAPLUS
DOCUMENT NUMBER: 137:353056
TITLE: Preparation of benzenesulfonylpiperazines as matrix
metalloproteinase inhibitors.
INVENTOR(S): Chung, Yong-Jun; Lee, Keyong-Ho; Kim, Youn-Chul; Park,
Ho-Jin
PATENT ASSIGNEE(S): Kolon Ind. Inc., S. Korea
SOURCE: PCT Int. Appl., 71 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002088115	A1	20021107	WO 2002-KR759	20020424
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG			
PRIORITY APPLN. INFO.:		KR 2001-22767	A	20010426
		KR 2001-77522	A	20011207
		KR 2002-14481	A	20020318
OTHER SOURCE(S):	MARPAT 137:353056			
GT				

OTHER SOURCE(S): MARPAT 137:353056
GI



AB Title compds. [I; n = 0-3; A = CO₂H, CONHOH, CH₂SH, CH₂OH; B = H, alkyl, NO₂, aryl, heteroaryl, pyrrolyl, halo, alkoxy, aryloxy, alkylamino, alkylthio, CONHR, NHCOR, NHCO₂R, NHCONHR, etc.; R = H, alkyl, aryl, heteroaryl, tetragonal to octagonal cyclic compd., alkyl substituted by a tetragonal to octagonal (hetero)cyclic compd.; Z = H, O, S, provided that when Z = O, S it takes a double bond; Y = H, alkyl, aryl, heteroaryl, alkyl substituted by a tetragonal to octagonal cyclic compd., alkyl substituted by a tetragonal to octagonal heterocycl, CONHR, NHCOR, NHCO₂R, NHCONHR, alkyl having a double or triple bond], were prep'd. Thus, Me 1-(4-methoxybenzenesulfonyl)-5-oxopiperazine-2-carboxylate (prepn. given) was stirred 5 h with aq. NH₂OH to give 45% 1-(4-methoxybenzenesulfonyl)-5-oxopiperazine-2-hydroxamic acid. This inhibited MMP-2 with IC₅₀ = 0.004 .μ.M. I are angiogenesis controlling materials that can inhibit overexpression of matrix **metalloproteinase** that decomp. protein constituents in extracellular matrix and basement membranes of connective tissues.

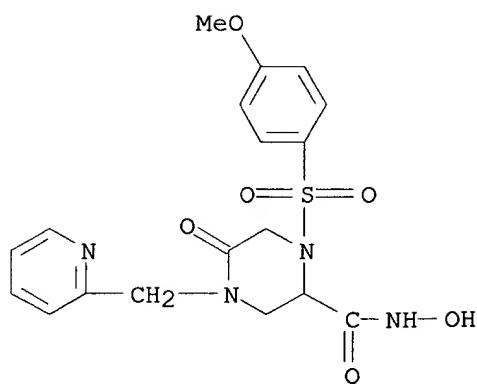
IT **474410-22-5P 474410-24-7P**

RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of benzenesulfonylpiperazines as matrix
metalloproteinase inhibitors)

RN 474410-22-5 CAPLUS

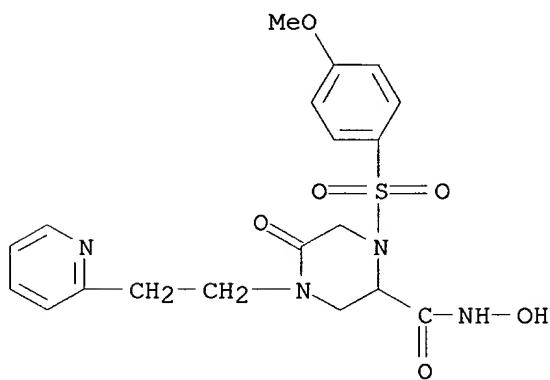
CN 2-Piperazinecarboxamide, N-hydroxy-1-[(4-methoxyphenyl)sulfonyl]-5-oxo-4-(2-pyridinylmethyl)-, monohydrochloride (9CI) (CA INDEX NAME)



HC1

RN 474410-24-7 CAPLUS

CN 2-Piperazinecarboxamide, N-hydroxy-1-[(4-methoxyphenyl)sulfonyl]-5-oxo-4-[2-(2-pyridinyl)ethyl]- (9CI) (CA INDEX NAME)

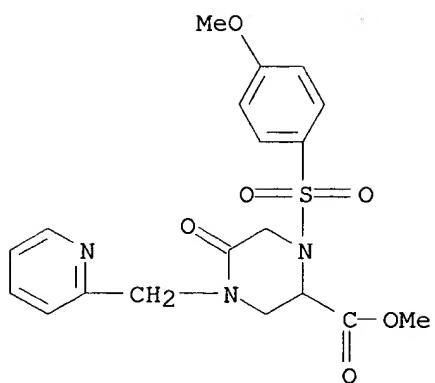


IT 474410-45-2P 474410-46-3P

RL: RCT (Reactant); SPN (Synthetic preparation); PREP (Preparation); RACT (Reactant or reagent)
(prepn. of benzenesulfonylpiperazines as matrix
metalloproteinase inhibitors)

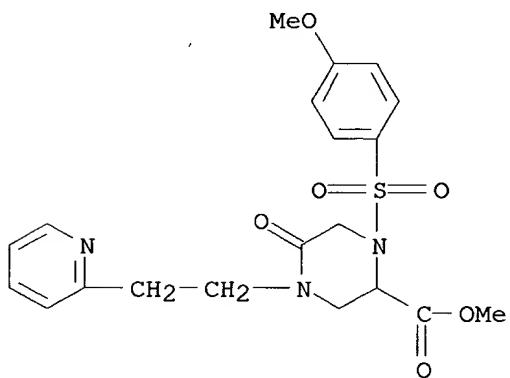
BN 474410-45-2 CAPTUS

CN 2-Piperinecarboxylic acid, 1-[(4-methoxyphenyl)sulfonyl]-5-oxo-4-(2-pyridinylmethyl)-, methyl ester (9CI) (CA INDEX NAME)



RN 474410-46-3 CAPLUS

CN 2-Piperazinecarboxylic acid, 1-[(4-methoxyphenyl)sulfonyl]-5-oxo-4-[2-(2-pyridinyl)ethyl]-, methyl ester (9CI) (CA INDEX NAME)



REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 3 OF 15 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2002:849440 CAPLUS

DOCUMENT NUMBER: 137:333179

TITLE: Use of HIV protease inhibitors to block cell migration and/or invasion, tissue infiltration, and edema, and therapeutic use

INVENTOR(S): Ensoli, Barbara

PATENT ASSIGNEE(S): Istituto Superiore di Sanita', Italy

SOURCE: PCT Int. Appl., 93 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002087583	A2	20021107	WO 2002-EP4303	20020418
WO 2002087583	A3	20021219		

W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM

RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG

PRIORITY APPLN. INFO.: IT 2001-RM210 A 20010418

AB The invention provides a method using HIV protease inhibitors for blocking the invasion of normal, neoplastic, inflammatory, or immune cells, tissue infiltration, and/or edema formation through inhibition or modulation of mols. and proteolytic enzymes (e.g. matrix **metalloproteinases**), for the therapy of diseases whose pathogenesis is related to the above processes, including tumors, non-neoplastic angioproliferative diseases, inflammatory diseases, or autoimmune diseases.

IT 150378-17-9, Indinavir 150378-17-9D, Indinavir, derivs.

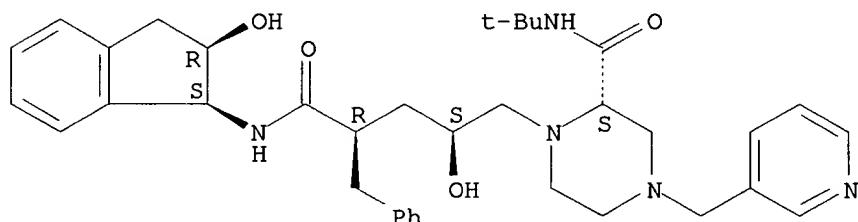
RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(HIV protease inhibitors to block cell migration and/or invasion, tissue infiltration, and edema, and therapeutic use)

RN 150378-17-9 CAPLUS

CN D-erythro-Pentonamide, 2,3,5-trideoxy-N-[(1S,2R)-2,3-dihydro-2-hydroxy-1H-inden-1-yl]-5-[(2S)-2-[(1,1-dimethylethyl)amino]carbonyl]-4-(3-pyridinylmethyl)-1-piperazinyl]-2-(phenylmethyl)- (9CI) (CA INDEX NAME)

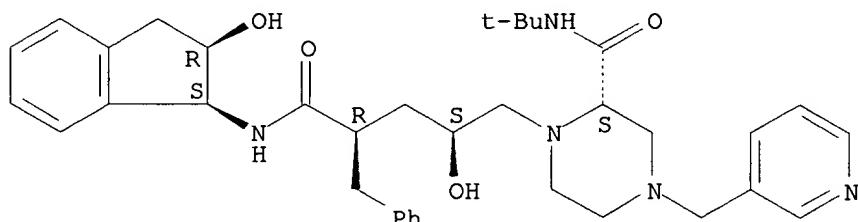
Absolute stereochemistry.



RN 150378-17-9 CAPLUS

CN D-erythro-Pentonamide, 2,3,5-trideoxy-N-[(1S,2R)-2,3-dihydro-2-hydroxy-1H-inden-1-yl]-5-[(2S)-2-[(1,1-dimethylethyl)amino]carbonyl]-4-(3-pyridinylmethyl)-1-piperazinyl]-2-(phenylmethyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L5 ANSWER 4 OF 15 CAPLUS COPYRIGHT 2003 ACS
 ACCESSION NUMBER: 2002:638144 CAPLUS
 DOCUMENT NUMBER: 137:163841
 TITLE: Methods for regulating levels of zinc, cadmium, and calcium in humans and for diagnosing, or screening for the risk of developing diseases associated with abnormal levels of cadmium, zinc and calcium in body fluids and tissues
 INVENTOR(S): Woods, Gordon L.
 PATENT ASSIGNEE(S): USA
 SOURCE: U.S. Pat. Appl. Publ., 19 pp., Cont.-in-part of U.S. Ser. No. 610,538, abandoned.
 CODEN: USXXCO
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 3
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
US 2002114848	A1	20020822	US 2001-989674	20011121
PRIORITY APPLN. INFO.:			US 1999-142926P	P 19990709
			US 2000-610538	B2 20000707

AB Methods and compns. are provided for decreasing PGE2:PGF2.alpha., regulating ratios of zinc:cadmium and regulating the concn. of zinc, calcium and zinc-contg. and PGE2-dependent matrix **metalloproteinases** in body fluids and tissues of a human. Elevated or otherwise unregulated levels of PGE2, zinc and calcium and elevated concns. of zinc-contg. and PGE2-dependent matrix **metalloproteinases** have been found to be assocd. with the development of certain diseases. Methods for the prevention of a variety of diseases are also disclosed.

IT 157810-81-6, Indinavir sulfate
 RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
 (zinc, cadmium, and calcium level regulation in humans, and use in disease diagnosis and prevention)

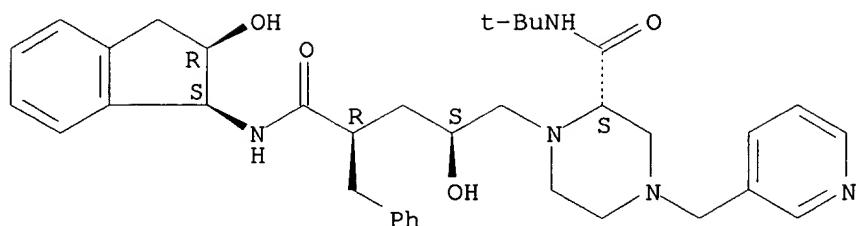
RN 157810-81-6 CAPLUS

CN D-erythro-Pentonamide, 2,3,5-trideoxy-N-[(1S,2R)-2,3-dihydro-2-hydroxy-1H-inden-1-yl]-5-[(2S)-2-[(1,1-dimethylethyl)amino]carbonyl]-4-(3-pyridinylmethyl)-1-piperazinyl]-2-(phenylmethyl)-, sulfate (1:1) (salt) (9CI) (CA INDEX NAME)

CM 1

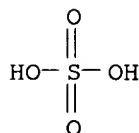
CRN 150378-17-9
 CMF C36 H47 N5 O4

Absolute stereochemistry.



CM 2

CRN 7664-93-9

CMF H₂ O₄ S

L5 ANSWER 5 OF 15 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2002:521462 CAPLUS

DOCUMENT NUMBER: 137:88442

TITLE: Incensole and furanogermacrens and compounds in treatment for inhibiting neoplastic lesions and microorganisms

INVENTOR(S): Shanahan-Pendergast, Elisabeth

PATENT ASSIGNEE(S): Ire.

SOURCE: PCT Int. Appl., 68 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2002053138	A2	20020711	WO 2002-IE1	20020102
WO 2002053138	A3	20020919		
W:	AE, AG, AT, AU, BB, BG, CA, CH, CN, CO, CU, CZ, LU, LV, MA, MD, UA, UG, US, VN, YU, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, SD, SL, SZ, UG, AT, BE, CH, CY, DE, ES, FI, ML, MR, NE, SN, TD, TG			

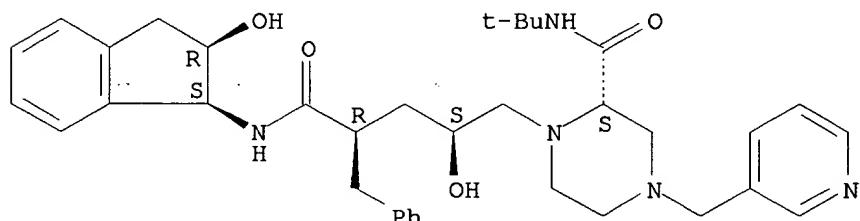
PRIORITY APPLN. INFO.: IE 2001-2 A 20010102

OTHER SOURCE(S): MARPAT 137:88442

AB The invention discloses the use of incensole and/or furanogermacrens, derivs. metabolites and precursors thereof in the treatment of neoplasia, particularly resistant neoplasia and immunodysregulatory disorders. These compds. can be administered alone or in combination with conventional chemotherapeutic, antiviral, antiparasite agents, radiation and/or surgery. Incensole and furanogermacren and their mixt. showed antitumor activity against various human carcinomas and melanomas and antimicrobial

activity against *Staphylococcus aureus* and *Enterococcus faecalis*.
 IT 150378-17-9, Indinavir
 RL: PAC (Pharmacological activity); THU (Therapeutic use); BIOL
 (Biological study); USES (Uses)
 (pharmaceutical formulation further contg.; incensole and
 furanogermacrens and compds. as antitumor and antimicrobial agents)
 RN 150378-17-9 CAPLUS
 CN D-erythro-Pentonamide, 2,3,5-trideoxy-N-[(1S,2R)-2,3-dihydro-2-hydroxy-1H-
 inden-1-yl]-5-[(2S)-2-[(1,1-dimethylethyl)amino]carbonyl]-4-(3-
 pyridinylmethyl)-1-piperazinyl]-2-(phenylmethyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L5 ANSWER 6 OF 15 CAPLUS COPYRIGHT 2003 ACS
 ACCESSION NUMBER: 2002:223957 CAPLUS
 DOCUMENT NUMBER: 137:88006
 TITLE: HIV protease inhibitors are potent anti-angiogenic molecules and promote regression of Kaposi sarcoma
 Sgadari, Cecilia; Barillari, Giovanni; Toschi, Elena;
 Carlei, Davide; Bacigalupo, Ilaria; Baccarini, Sara;
 Palladino, Clelia; Leone, Patrizia; Bugarini, Roberto;
 Malavasi, Laura; Cafaro, Aurelio; Falchi, Mario;
 Valdembri, Donatella; Rezza, S, Giovanni; Bussolino, Federico; Monini, Paolo; Ensoli, Barbara
 CORPORATE SOURCE: Laboratory of Virology, Istituto Superiore di Sanita, Rome, Italy
 SOURCE: Nature Medicine (New York, NY, United States) (2002), 8(3), 225-232
 CODEN: NAMEFI; ISSN: 1078-8956
 PUBLISHER: Nature America Inc.
 DOCUMENT TYPE: Journal
 LANGUAGE: English
 AB Treatment with HIV-1 protease inhibitors (PI) is assocd. with a reduced incidence or regression of Kaposi sarcoma (KS). Here we show that systemic administration of the PIs indinavir or saquinavir to nude mice blocks the development and induces regression of angioproliferative KS-like lesions promoted by primary human KS cells, basic fibroblast growth factor (bFGF), or bFGF and vascular endothelial growth factor (VEGF) combined. These PIs also block bFGF or VEGF-induced angiogenesis in the chorioallantoic membrane assay with a potency similar to paclitaxel (Taxol). These effects are mediated by the inhibition of endothelial- and KS-cell invasion and of matrix metalloproteinase-2 proteolytic activation by PIs at concns. present in plasma of treated individuals. As PIs also inhibit the in vivo growth and invasion of an angiogenic tumor-cell line, these data indicate that PIs are potent anti-angiogenic and anti-tumor mols. that might be used in treating non-HIV KS and in

other HIV-assocd. tumors.

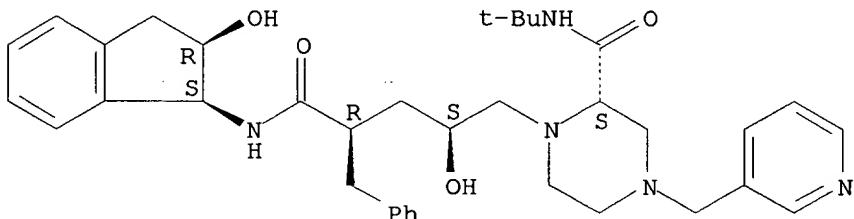
IT 150378-17-9, Indinavir

RL: DMA (Drug mechanism of action); PAC (Pharmacological activity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(HIV protease inhibitors are potent anti-angiogenic mols. and promote regression of Kaposi sarcoma)

RN 150378-17-9 CAPLUS

CN D-erythro-Pentonamide, 2,3,5-trideoxy-N-[(1S,2R)-2,3-dihydro-2-hydroxy-1H-inden-1-yl]-5-[(2S)-2-[(1,1-dimethylethyl)amino]carbonyl]-4-(3-pyridinylmethyl)-1-piperazinyl]-2-(phenylmethyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT: 50 THERE ARE 50 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 7 OF 15 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2002:46330 CAPLUS

DOCUMENT NUMBER: 137:134505

TITLE: Synergistic antiadipogenic effects of HIV type 1 protease inhibitors with tumor necrosis factor .alpha.: Suppression of extracellular insulin action mediated by extracellular matrix-degrading proteases

AUTHOR(S): Mondal, Debasis; Larussa, Vincent F.; Agrawal, Krishna C.

CORPORATE SOURCE: Department of Pharmacology, Tulane University School of Medicine, New Orleans, LA, 70112, USA

SOURCE: AIDS Research and Human Retroviruses (2001), 17(17), 1569-1584

CODEN: ARHRE7; ISSN: 0889-2229

PUBLISHER: Mary Ann Liebert, Inc.

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Long-term use of HIV-1 protease inhibitors (PIs) is assocd. with a lipodystrophy syndrome. To delineate the assocd. mechanisms, adipogenesis was detd. in 3T3-L1 cells in the presence or absence of either indinavir (2-50 .mu.g/mL) or ritonavir (0.4-10 .mu.g/mL). A concn.-dependent decrease in both lipid (4-59%) and triglyceride (11-49%) levels was seen after 10 days of exposure. Simultaneous treatment with TNF-.alpha. showed a synergistic suppression in lipid levels by 45-95% at 10 U/mL and almost complete suppression at 100 U/mL. The effect of PIs on insulin-induced lipogenesis was monitored by [14C]glucose incorporation into lipids, which was suppressed by 21-86% in a concn.-dependent manner.

Insulin-sensitizing agent, troglitazone (80 and 400 nM), effectively blocked the PI-mediated adipogenic suppression. Preadipocyte factor 1 gene (pref-1) expression, as monitored by RT-PCR, was down-regulated (4- to 6-fold) within 48 h after insulin stimulation; however, a smaller

decrease (1.2- to 1.8-fold) was obsd. in PI-exposed cells. The decrease in proteolytic activity of matrix metalloproteases (MMP-2 and MMP-9) during adipogenesis was reversed on exposure to the PIs. Similarly, the plasminolytic activity was increased and plasminogen activator inhibitor (PAI) activity was decreased in supernatants from PI-treated cells. The insulin-mediated induction (3- to 4-fold) of PAI-1 and PAI-2 message was suppressed on exposure to PIs, which was reversed by troglitazone treatment. Thus, the HIV-1 PIs may suppress adipogenesis by disrupting the concerted actions of host proteases that regulate ECM integrity required for initiation of differentiation.

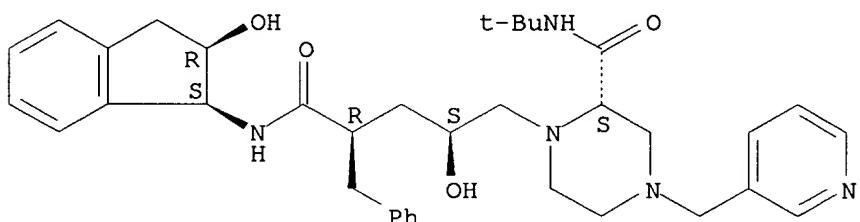
IT 150378-17-9, Indinavir

RL: ADV (Adverse effect, including toxicity); THU (Therapeutic use); BIOL (Biological study); USES (Uses)
(synergistic antiadipogenic effects of HIV type 1 proteinase inhibitors with TNF-.alpha. with suppression of extracellular insulin action mediated by extracellular matrix-degrading proteinases)

RN 150378-17-9 CAPLUS

CN D-erythro-Pentonamide, 2,3,5-trideoxy-N-[(1S,2R)-2,3-dihydro-2-hydroxy-1H-inden-1-yl]-5-[(2S)-2-[[[(1,1-dimethylethyl)amino]carbonyl]-4-(3-pyridinylmethyl)-1-piperazinyl]-2-(phenylmethyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT:

67

THERE ARE 67 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 8 OF 15 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2001:749723 CAPLUS

DOCUMENT NUMBER: 136:31301

TITLE: Three-Dimensional Quantitative Structure-Activity Relationship (3D-QSAR) Models for a Novel Class of Piperazine-Based Stromelysin-1 (MMP-3) Inhibitors: Applying a "Divide and Conquer" Strategy

AUTHOR(S): Amin, Elizabeth Ambrose; Welsh, William J.

CORPORATE SOURCE: Department of Chemistry & Biochemistry, University of Missouri-St. Louis, St. Louis, MO, 63121, USA

SOURCE: Journal of Medicinal Chemistry (2001), 44(23), 3849-3855

CODEN: JMCMAR; ISSN: 0022-2623

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal

LANGUAGE: English

AB Three-dimensional quant. structure-activity relationship (3D-QSAR) models have been obtained using comparative mol. field anal. (CoMFA) for a novel series of piperazine-based matrix metalloproteinase inhibitors (MMPIs). The crystal structure of stromelysin-1 (MMP-3) was used to identify regions of the enzyme and inhibitors where steric and

electrostatic effects correlate strongly with biol. activity. A training set composed of a subset of inhibitors (#10-35), which differed only with regards to the substituent (n-alkyl, amide, carbamide and sulfonamide) on the piperazine distal nitrogen, yielded the most predictive CoMFA model, with r^2 values of 0.592 (cross-validated) and 0.989 (conventional); this model was further validated using test compds. from two inhibitor subsets. Investigation of various ligand conformations, inhibitor subsets, alignment schemes and partial charge formalisms was required to obtain satisfactory models. The greatest success was achieved by incorporating inertial alignment together with manual adjustment of the enzyme-docked inhibitors to ensure complementarity between the inhibitors' substituent conformations and the structural characteristics of the MMP-3 S1-S2' binding pockets. Key insights into the structure-activity relationship (SAR) obtained from this anal. for this inhibitor set are in agreement with exptl. obsd. data on stromelysin-1 biol. activity and binding-site topol. In particular, the present study sheds new light on the steric and electrostatic requirements for ligand binding to the partly solvent-exposed S1-S2' area.

IT

262420-38-2

RL: BSU (Biological study, unclassified); PAC (Pharmacological activity); PRP (Properties); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

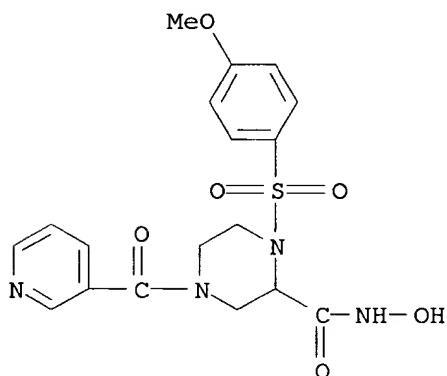
(three-dimensional quant. structure-activity relationship (3D-QSAR) models for a novel class of piperazine-based stromelysin-1 (MMP-3) inhibitors)

RN

262420-38-2 CAPLUS

CN

2-Piperazinecarboxamide, N-hydroxy-1-[(4-methoxyphenyl)sulfonyl]-4-(3-pyridinylcarbonyl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT:

20

THERE ARE 20 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 9 OF 15 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2001:564870 CAPLUS

DOCUMENT NUMBER: 135:132456

TITLE: Treatment of neuropsychiatric diseases with protease and neuraminidase inhibitors, and screening method

INVENTOR(S): Vawter, Marquis P.; Freed, William J.

PATENT ASSIGNEE(S): Department of Health and Human Services, USA

SOURCE: PCT Int. Appl., 73 pp.

CODEN: PIXXD2

DOCUMENT TYPE:

Patent

LANGUAGE:

English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001054729	A1	20010802	WO 2001-US2417	20010125
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				
AU 2001031137	A5	20010807	AU 2001-31137	20010125
PRIORITY APPLN. INFO.:			US 2000-177971P	P 200000125
			WO 2001-US2417	W 20010125

AB The invention provides a method of treating a neuropsychiatric disease characterized by an abnormally elevated level of a fragment of an isoform of a neural cell adhesion mol. (N-CAM) in the brain or cerebrospinal fluid of an affected human subject, comprising administering a therapeutically effective amt. of at least one compd. selected from protease inhibitors and neuraminidase inhibitors, whereby administering the compd. to the subject treats the human subject. The invention further provides a method of monitoring the efficacy of treatment with the method of the present invention. Moreover, the invention provides a method of screening for compds. effective in treating neuropsychiatric disease assocd. with an abnormally elevated level of a fragment of a N-CAM in the cerebrospinal fluid of an affected human subject. Further provided are fragments of an isoform of N-CAM in the cerebrospinal fluid of human subjects.

IT 150378-17-9, Indinavir

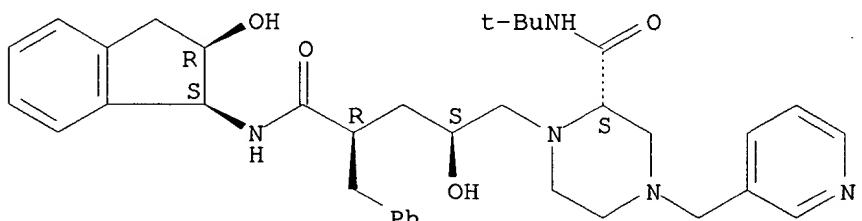
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(protease and neuraminidase inhibitors for treatment of neuropsychiatric diseases, and screening method)

RN 150378-17-9 CAPLUS

CN D-erythro-Pentonamide, 2,3,5-trideoxy-N-[(1S,2R)-2,3-dihydro-2-hydroxy-1H-inden-1-yl]-5-[(2S)-2-[[1,1-dimethylethyl]amino]carbonyl]-4-(3-pyridinylmethyl)-1-piperazinyl]-2-(phenylmethyl)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



REFERENCE COUNT:

9

THERE ARE 9 CITED REFERENCES AVAILABLE FOR THIS

RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 10 OF 15 CAPLUS COPYRIGHT 2003 ACS
 ACCESSION NUMBER: 2001:338762 CAPLUS
 DOCUMENT NUMBER: 134:362292
 TITLE: Methods of determining individual hypersensitivity to a pharmaceutical agent from gene expression profile
 INVENTOR(S): Farr, Spencer
 PATENT ASSIGNEE(S): Phase-1 Molecular Toxicology, USA
 SOURCE: PCT Int. Appl., 222 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001032928	A2	20010510	WO 2000-US30474	20001103
WO 2001032928	A3	20020725		
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG				

PRIORITY APPLN. INFO.: US 1999-165398P P 19991105
 US 2000-196571P P 20000411

AB The invention discloses methods, gene databases, gene arrays, protein arrays, and devices that may be used to det. the hypersensitivity of individuals to a given agent, such as drug or other chem., in order to prevent toxic side effects. In one embodiment, methods of identifying hypersensitivity in a subject by obtaining a gene expression profile of multiple genes assocd. with hypersensitivity of the subject suspected to be hypersensitive, and identifying in the gene expression profile of the subject a pattern of gene expression of the genes assocd. with hypersensitivity are disclosed. The gene expression profile of the subject may be compared with the gene expression profile of a normal individual and a hypersensitive individual. The gene expression profile of the subject that is obtained may comprise a profile of levels of mRNA or cDNA. The gene expression profile may be obtained by using an array of nucleic acid probes for the plurality of genes assocd. with hypersensitivity. The expression of the genes predetd. to be assocd. with hypersensitivity is directly related to prevention or repair of toxic damage at the tissue, organ or system level. Gene databases arrays and app. useful for identifying hypersensitivity in a subject are also disclosed.

IT 150378-17-9, Indinavir

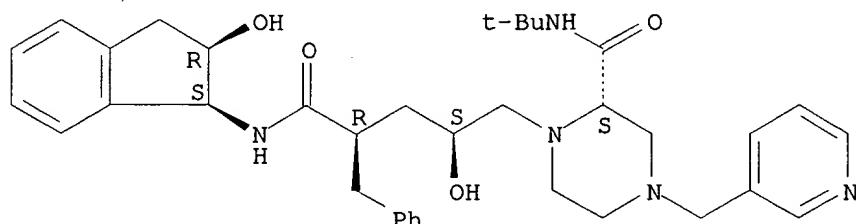
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); BIOL (Biological study)
 (methods of detg. individual hypersensitivity to a pharmaceutical agent from gene expression profile)

RN 150378-17-9 CAPLUS

CN D-erythro-Pentonamide, 2,3,5-trideoxy-N-[(1S,2R)-2,3-dihydro-2-hydroxy-1H-

inden-1-yl]-5-[(2S)-2-[[[(1,1-dimethylethyl)amino]carbonyl]-4-(3-pyridinylmethyl)-1-piperazinyl]-2-(phenylmethyl)-(9CI) (CA INDEX NAME)

Absolute stereochemistry.



L5 ANSWER 11 OF 15 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2001:247178 CAPLUS

DOCUMENT NUMBER: 134:275776

TITLE: Method using a geranylgeranyl-protein transferase inhibitor for preventing osteoporosis, pharmaceutical compositions, and compound preparation

INVENTOR(S): Reszka, Alfred A.; Rodan, Gideon A.

PATENT ASSIGNEE(S): Merck & Co., Inc., USA

SOURCE: PCT Int. Appl., 210 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001022963	A1	20010405	WO 2000-US26357	20000925
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW:	GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG			

PRIORITY APPLN. INFO.: US 1999-156234P P 19990927

OTHER SOURCE(S): MARPAT 134:275776

AB A method for preventing or inhibiting bone resorption in a mammal comprises administering to a mammal in need thereof a therapeutically effective amt. of an inhibitor of geranylgeranyl-protein transferase type I.

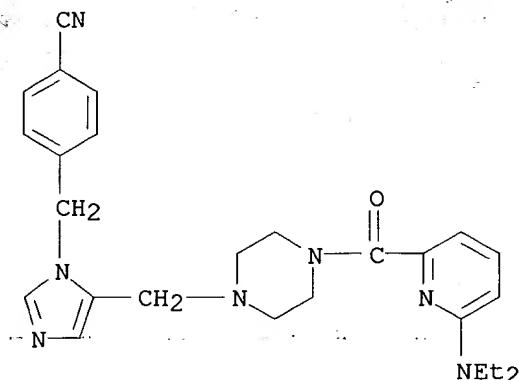
IT 290819-82-8P

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(geranylgeranyl-protein transferase inhibitor for preventing bone resorption, pharmaceutical compns., and compd. prepn.)

RN 290819-82-8 CAPLUS

CN Piperazine, 1-[(1-[(4-cyanophenyl)methyl]-1H-imidazol-5-yl)methyl]-4-[(6-

(diethylamino)-2-pyridinyl]carbonyl]-, trihydrochloride (9CI) (CA INDEX NAME).



● 3 HCl

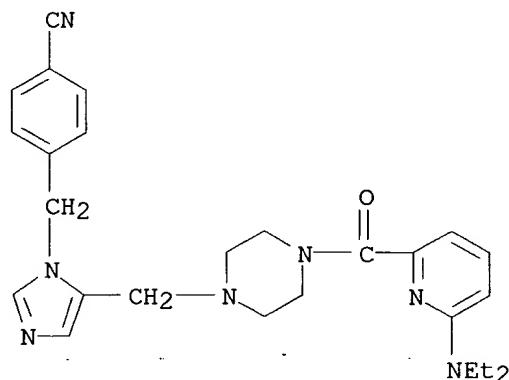
IT 290819-51-1

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(geranylgeranyl-protein transferase inhibitor for preventing bone resorption, pharmaceutical compns., and compd. prepns.)

RN 290819-51-1 CAPLUS

CN Piperazine, 1-[(1-[(4-cyanophenyl)methyl]-1H-imidazol-5-yl)methyl]-4-[(6-diethylamino)-2-pyridinyl]carbonyl]- (9CI) (CA INDEX NAME)



REFERENCE COUNT:

4

THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 12 OF 15 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2001:50495 CAPLUS

DOCUMENT NUMBER: 134:95488

TITLE: Cadmium containing compositions for prevention and

INVENTOR(S): treatment of prostate cancer
 Woods, Gordon L.
 PATENT ASSIGNEE(S): Cancer2 Inc., USA
 SOURCE: PCT Int. Appl., 42 pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: English
 FAMILY ACC. NUM. COUNT: 3
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2001003708	A1	20010118	WO 2000-US18580	20000707
W: CA RW: AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE				
EP 1200104	A1	20020502	EP 2000-947094	20000707
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, FI, CY				
US 2002148049	A1	20021017	US 2002-38035	20020102
PRIORITY APPLN. INFO.:			US 1999-142926P	P 19990709
			EP 1999-113269	A 19990708
			WO 2000-US18580	W 20000707

AB Methods and compns. are provided for decreasing or regulating ratios of zinc:cadmium and PGE2:PGF2.alpha. and regulating the concn. of zinc-contg. and PGE2-dependent matrix **metalloproteinases** in body fluids and tissues of a mammal, comprising administering to the mammal an amt. of a pharmaceutically acceptable and bioavailable cadmium salt. Elevated or fluctuating levels of PGE2 and zinc and elevated concns. of zinc-contg. and PGE2-dependent matrix **metalloproteinases** have been found to be assocd. with the development of certain diseases, e.g. prostate cancer, diabetes, and multiple sclerosis. Ejaculates from horse stallions showed that when the concn. of cadmium is increased, the sperm motility decreases. Motility of the sperm correlates to sperm viability which is an indicator of the proliferation environment of the stallion's prostate glands. Higher cadmium values from semen decreases the proliferation of prostate cells and replication of viruses within the prostate environment. Thus elevating the cadmium concn. in man's prostate gland will decrease man's incidence of prostate cancer, decreases his fertility, and protects against viral infections and age-onset diseases.

IT 157810-81-6, Indinavir sulfate

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); THU (Therapeutic use); BIOL (Biological study); USES (Uses)

(cadmium contg. compns. for prevention and treatment of prostate cancer)

RN 157810-81-6 CAPLUS

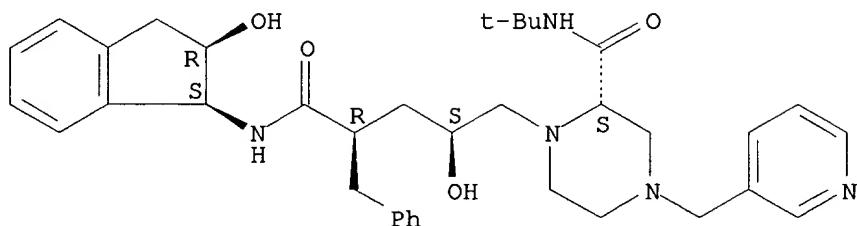
CN D-erythro-Pentonamide, 2,3,5-trideoxy-N-[(1S,2R)-2,3-dihydro-2-hydroxy-1H-inden-1-yl]-5-[(2S)-2-[(1,1-dimethylethyl)amino]carbonyl]-4-(3-pyridinylmethyl)-1-piperazinyl]-2-(phenylmethyl)-, sulfate (1:1) (salt) (9CI) (CA INDEX NAME)

CM 1

CRN 150378-17-9

CMF C36 H47 N5 O4

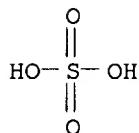
Absolute stereochemistry.



CM 2

CRN 7664-93-9

CMF H2 O4 S



REFERENCE COUNT: 6 THERE ARE 6 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 13 OF 15 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 2000:44069 CAPLUS

DOCUMENT NUMBER: 132:251126

TITLE: Design and Synthesis of Piperazine-Based Matrix **Metalloproteinase** Inhibitors

AUTHOR(S): Cheng, Menyan; De, Biswanath; Pikul, Stanislaw; Almstead, Neil G.; Natchus, Michael G.; Anastasio, Melanie V.; McPhail, Sara J.; Snider, Catherine E.; Taiwo, Yetunde O.; Chen, Longyin; Dunaway, C. Michelle; Gu, Fei; Dowty, Martin E.; Mieling, Glen E.; Janusz, Michael J.; Wang-Weigand, Sherry

CORPORATE SOURCE: Health Care Research Center, Procter and Gamble Pharmaceuticals, Mason, OH, 45040, USA

SOURCE: Journal of Medicinal Chemistry (2000), 43(3), 369-380
CODEN: JMCMAR; ISSN: 0022-2623

PUBLISHER: American Chemical Society

DOCUMENT TYPE: Journal

LANGUAGE: English

AB A new generation of cyclic matrix **metalloproteinase** (MMP) inhibitors derived from dl-piperazinecarboxylic acid was described. The design involves: incorporation of hydroxamic acid as the bidentate chelating agent for catalytic Zn²⁺, placement of a sulfonamide group at the 1N-position of the piperazine ring to fill the S1' pocket of the enzyme, and finally attachment of diverse functional groups at the 4N-position to optimize potency and peroral absorption. A unique combination of all three elements produced 3-[(hydroxyamino)carbonyl]-4-[(4-methoxyphenyl)sulfonyl]-1-piperazinecarboxylic acid phenylmethyl ester with high affinity for MMP-1, MMP-3, MMP-9, and MMP-13 (24, 18, 1.9, and

1.3 nM, resp.). X-ray crystallog. data obtained for MMP-3 co-crystd. with 3-[(hydroxyamino)carbonyl]-4-[(4-methoxyphenyl)sulfonyl]-1-piperazinecarboxylic acid phenylmethyl ester gave detailed information on key binding interactions defining an overall scaffold geometry for piperazine-based MMP inhibitors.

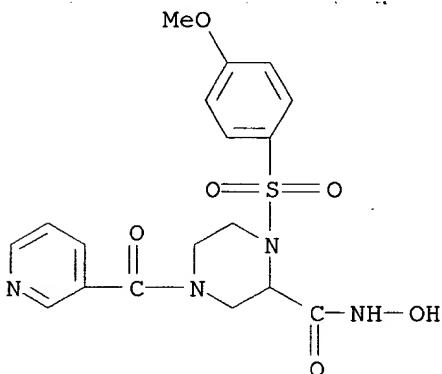
IT **262420-38-2P**

RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); BIOL (Biological study); PREP (Preparation)

(prepn. and activity of N-hydroxy(phenylsulfonyl)piperazinecarboxamides as matrix **metalloproteinase** inhibitors)

RN 262420-38-2 CAPLUS

CN 2-Piperazinecarboxamide, N-hydroxy-1-[(4-methoxyphenyl)sulfonyl]-4-(3-pyridinylcarbonyl)- (9CI) (CA INDEX NAME)



REFERENCE COUNT: 26 THERE ARE 26 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L5 ANSWER 14 OF 15 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1998:498326 CAPLUS

DOCUMENT NUMBER: 129:148991

TITLE: Preparation of N-sulfamoylpiperidine-2-hydroxamic acids and analogs as **metalloproteinase** inhibitors

INVENTOR(S): Broka, Chris Allen; Campbell, Jeffrey Allen; Castelhano, Arlindo Lucas; Chen, Jian Jeffrey; Hendricks, Robert Than; Melnick, Michael Joseph; Walker, Keith Adrian Murray

PATENT ASSIGNEE(S): F. Hoffmann-La Roche A.-G., Switz.; Agouron Pharmaceuticals, Inc.

SOURCE: Ger. Offen., 84 pp.

CODEN: GWXXBX

DOCUMENT TYPE: Patent

LANGUAGE: German

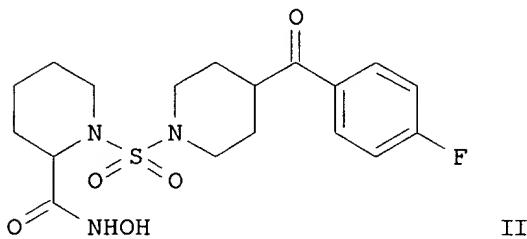
FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
DE 19802350	A1	19980730	DE 1998-19802350	19980122
WO 9832748	A1	19980730	WO 1998-EP180	19980114

W: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE,
 DK, EE, ES, FI, GB, GE, GH, GM, GW, HU, ID, IL, IS, JP, KE, KG,
 KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX,
 NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT,
 UA, UG, UZ, VN, YU, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
 RW: GH, GM, KE, LS, MW, SD, SZ, UG, ZW, AT, BE, CH, DE, DK, ES, FI,
 FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, BF, BJ, CF, CG, CI, CM,
 GA, GN, ML, MR, NE, SN, TD, TG
 AU 9866140 A1 19980818 AU 1998-66140 19980114
 AU 730127 B2 20010222
 EP 958287 A1 19991124 EP 1998-907943 19980114
 EP 958287 B1 20020911
 R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
 IE, SI, LT, LV, FI, RO
 BR 9807508 A 20000321 BR 1998-7508 19980114
 NZ 336625 A 20010427 NZ 1998-336625 19980114
 JP 2001523222 T2 20011120 JP 1998-531537 19980114
 AT 223909 E 20020915 AT 1998-907943 19980114
 CN 1093125 B 20021023 CN 1998-803233 19980114
 ZA 9800376 A 19980723 ZA 1998-376 19980116
 IT 1298163 B1 19991220 IT 1998-MI91 19980120
 FR 2758559 A1 19980724 FR 1998-601 19980121
 GB 2321641 A1 19980805 GB 1998-1393 19980122
 GB 2321641 B2 20010401
 ES 2136037 A1 19991101 ES 1998-113 19980122
 ES 2136037 B1 20001116
 NO 9903587 A 19990922 NO 1999-3587 19990722
 MX 9906822 A 20000131 MX 1999-6822 19990722
 PRIORITY APPLN. INFO.: US 1997-36714P P 19970123
 US 1997-62209P P 19971016
 WO 1998-EP180 W 19980114

OTHER SOURCE(S): MARPAT 129:148991
GI



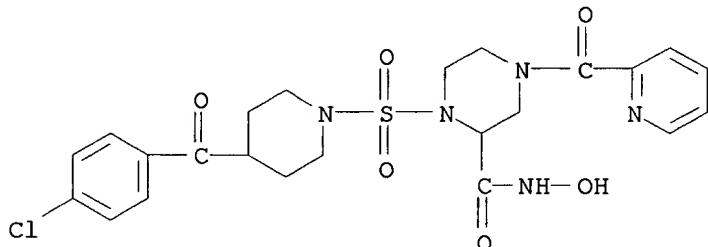
AB R10COCR1R2NR3SO2NR20R21 [I; R1-R3 = H, (CO-interrupted) alkyl,
 heterocycl(alkyl), (hetero)aryl(alkyl), etc.; R1R2, R1R3, R2R3 = atoms
 to complete a ring; R10 = NR11OR12; R11, R12 = H or (ar)alkyl; R20, R21 = H,
 alkyl, (hetero)aryl[alk(en)yl], etc.; NR20R21heterocycl] were prep'd.
 Thus, (R)-1-[4-(4-chlorobenzoyl)piperidine-1-sulfonyl]piperidine-2-
 carboxylic acid was amidated by H2NOCMe3 and the product deprotected to
 give title compd. (R)-II. Data for biol. activity of I were given.

IT 210915-42-7P 210916-04-4P
 RL: BAC (Biological activity or effector, except adverse); BSU (Biological
 study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use);
 BIOL (Biological study); PREP (Preparation); USES (Uses)

(prepn. of N-sulfamoylpiperidine-2-hydroxamic acids and analogs as metalloproteinase inhibitors)

RN 210915-42-7 CAPLUS

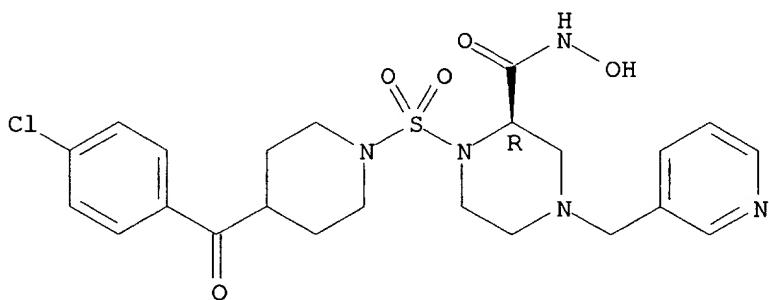
CN 2-Piperazinecarboxamide, 1-[[4-(4-chlorobenzoyl)-1-piperidinyl]sulfonyl]-N-hydroxy-4-(2-pyridinylcarbonyl)- (9CI) (CA INDEX NAME)



RN 210916-04-4 CAPLUS

CN 2-Piperazinecarboxamide, 1-[[4-(4-chlorobenzoyl)-1-piperidinyl]sulfonyl]-N-hydroxy-4-(3-pyridinylmethyl)-, (2R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



L5 ANSWER 15 OF 15 CAPLUS COPYRIGHT 2003 ACS

ACCESSION NUMBER: 1998:424232 CAPLUS

DOCUMENT NUMBER: 129:95510

TITLE: Preparation of 2-piperazinecarboxamides as inhibitors of MMP or TNF

INVENTOR(S): Neya, Masahiro; Yamazaki, Hitoshi; Kayakiri, Natsuko; Sato, Kentaro; Oku, Teruo

PATENT ASSIGNEE(S): Fujisawa Pharmaceutical Co., Ltd., Japan; Neya, Masahiro; Yamazaki, Hitoshi; Kayakiri, Natsuko; Sato, Kentaro; Oku, Teruo

SOURCE: PCT Int. Appl., 199 pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: English

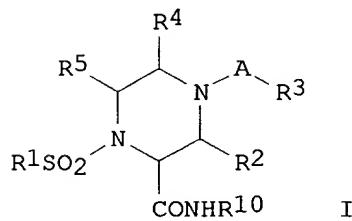
FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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WO 9827069	A1 19980625	WO 1997-JP4613	19971215
W: AU, CA, CN, HU, IL, JP, KR, MX, US, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
RW: AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE			
ZA 9711284	A 19980623	ZA 1997-11284	19971215
AU 9854122	A1 19980715	AU 1998-54122	19971215
EP 948489	A1 19991013	EP 1997-947944	19971215
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, PT, IE, FI			
JP 2001506257	T2 20010515	JP 1998-527536	19971215
KR 2000057595	A 20000925	KR 1999-705374	19990615
US 6333324	B1 20011225	US 1999-319928	19990726
US 2002128270	A1 20020912	US 2001-982869	20011022
US 6489324	B2 20021203		
PRIORITY APPLN. INFO.:			
		AU 1996-4249	A 19961217
		AU 1997-7156	A 19970603
		AU 1997-8568	A 19970814
		WO 1997-JP4613	W 19971215
		US 1999-319928	A3 19990726

OTHER SOURCE(S): MARPAT 129:95510
GI



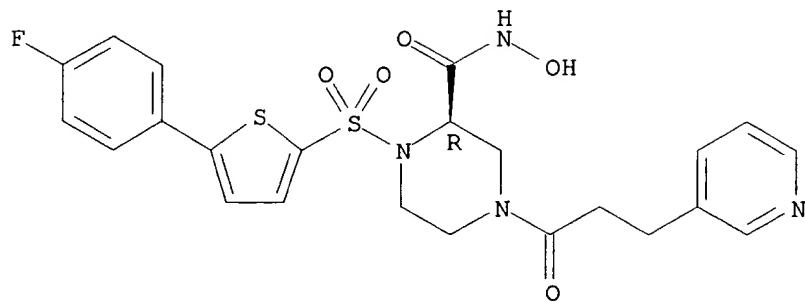
AB The title compds. [I; A = SO₂, C(O); R₁ = (un)substituted aryl, heterocyclic, lower alkyl, lower alkenyl; R₂ = H, (un)substituted lower alkyl, aryl, heterocyclic; R₃ = (un)substituted lower alkyl, lower alkoxy, aryloxy, etc.; R₄ = H, (un)substituted lower alkyl, aryl, heterocyclic; R₅ = H, (un)substituted lower alkyl, aryl, heterocyclic; R₁₀ = OH, protected OH] and their pharmaceutically acceptable salts, useful for prophylactic and therapeutic treatment of MMP- or TNF. α -mediated diseases, were prepd. Thus, treatment of a soln. of (2R)-1-(4-nitrobenzenesulfonyl)-4-methanesulfonylpiperazine-2-[N-(2-tetrahydropyran-2-yl)]carboxamide in MeOH with 10% HCl-MeOH afforded (2R)-I [R₁ = 4-O₂NC₆H₄SO₂; R₂ = R₄ = R₅ = H; R₃ = Me; R₁₀ = OH] which showed 95.3% inhibition of collagenase activity at 1x10⁻⁶ M.

IT 209590-20-5P 209590-22-7P 209591-56-0P
209591-57-1P
RL: BAC (Biological activity or effector, except adverse); BSU (Biological study, unclassified); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(prepn. of 2-piperazinecarboxamides as inhibitors of MMP or TNF)

RN 209590-20-5 CAPLUS

CN 2-Piperazinecarboxamide, 1-[[5-(4-fluorophenyl)-2-thienyl]sulfonyl]-N-hydroxy-4-[1-oxo-3-(3-pyridinyl)propyl]-, monohydrochloride, (2R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.



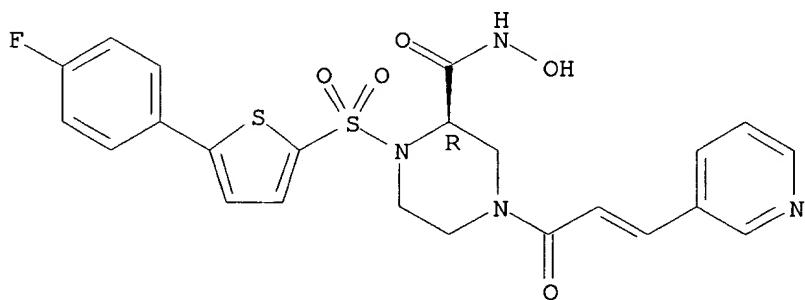
● HCl

RN 209590-22-7 CAPLUS

CN 2-Piperazinecarboxamide, 1-[[5-(4-fluorophenyl)-2-thienyl]sulfonyl]-N-hydroxy-4-[1-oxo-3-(3-pyridinyl)-2-propenyl]-, monohydrochloride, (2R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry unknown.

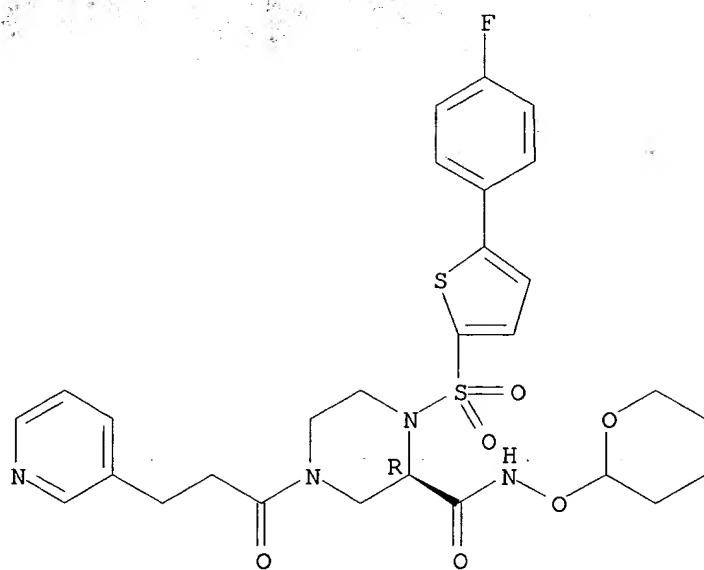


● HCl

RN 209591-56-0 CAPLUS

CN 2-Piperazinecarboxamide, 1-[[5-(4-fluorophenyl)-2-thienyl]sulfonyl]-4-[1-oxo-3-(3-pyridinyl)propyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]-, (2R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

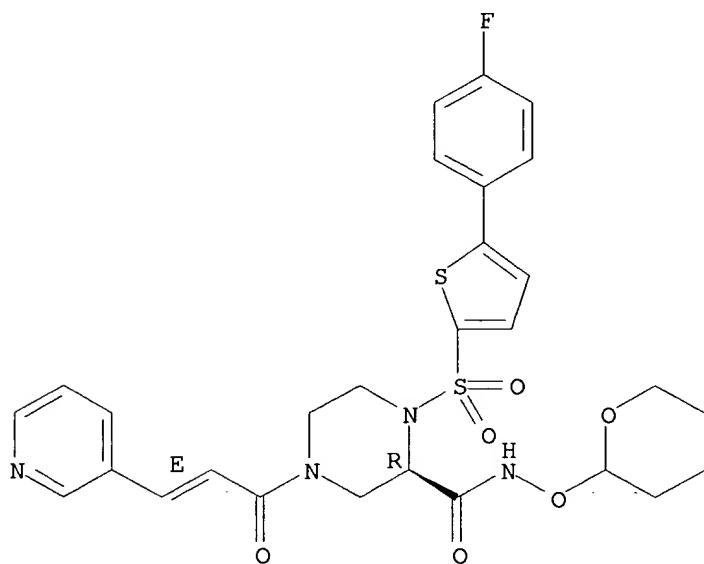


RN 209591-57-1 CAPLUS

CN 2-Piperazinecarboxamide, 1-[[5-(4-fluorophenyl)-2-thienyl]sulfonyl]-4-[(2E)-1-oxo-3-(3-pyridinyl)-2-propenyl]-N-[(tetrahydro-2H-pyran-2-yl)oxy]-, (2R)- (9CI) (CA INDEX NAME)

Absolute stereochemistry.

Double bond geometry as shown.



REFERENCE COUNT:

3

THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

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Page 33

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FULL ESTIMATED COST	70.60	218.96
DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)	SINCE FILE ENTRY	TOTAL SESSION
CA SUBSCRIBER PRICE	-9.77	-9.77

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